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- (54) **1-(4-SULFAMYLARYL)-3-SUBSTITUTED-5-ARYL-2-PYRAZOLINES AND INHIBITORS OF CYCLOOXYGENASE-2**
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- (51) **Int. Cl.<sup>7</sup>** ..... C07D 231/06; C07D 401/04; C07D 405/04; A61K 31/402; A61K 31/4439
- (52) **U.S. Cl.** ..... 514/341; 514/403; 546/275.4; 548/364.4; 548/364.7; 548/365.7; 548/379.1; 548/379.7
- (58) **Field of Search** ..... 548/365.7, 379.1, 548/379.7, 364.4, 364.7; 546/275.4; 514/341, 403

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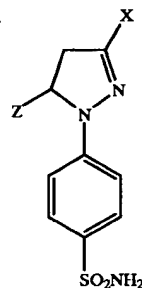
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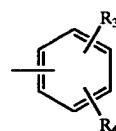
*Primary Examiner*—Fiona T. Powers(74) *Attorney, Agent, or Firm*—Drinker Biddle & Reath LLP(57) **ABSTRACT**

Compounds of the formula



(I)

wherein:

X is selected from the group consisting of trihalomethyl, C<sub>1</sub>-C<sub>6</sub> alkyl, and a group of formula II:

(II)

wherein:

R<sub>3</sub> and R<sub>4</sub> are independently selected from the group consisting of hydrogen; halogen; hydroxyl; nitro; C<sub>1</sub>-C<sub>6</sub> alkyl;C<sub>1</sub>-C<sub>6</sub> alkoxy; carboxy; C<sub>1</sub>-C<sub>6</sub> trihaloalkyl; and cyano;

Z is selected from the group consisting of substituted and unsubstituted aryl; or a pharmaceutically acceptable salt thereof. The compounds are inhibitors of cyclooxygenase-2 activity. They are useful for treating cyclooxygenase-mediated disorders, including, for example, inflammation, neoplastic disorders and angiogenesis-mediated disorders.